## **AMENDMENTS TO THE CLAIMS**

This listing of claims will replace all prior versions, and listings of claims in the application:

Claims 1-26 (canceled)

Claim 27 (currently amended): A method for enhancing efficacy of a chemotherapeutic agent for a cancer cell, said method comprising administering systemically to a subject in need thereof an effective amount of hyaluronan and said chemotherapeutic agent, wherein the hyaluronan has a molecular weight between 400,000 and 900,000 modal molecular weight of 890,000 Da.

Claims 28-29 (canceled)

Claim 30 (currently amended): The method according to Claim [[28]] <u>27</u>, wherein the hyaluronan has a molecular weight of 890,000 Da.

Claim 31 (canceled)

Claim 32 (currently amended): The method according to Claim [[28]] <u>27</u>, wherein the chemotherapeutic agent is selected from the group consisting of methotrexate, paclitaxel, 5-fluorouracil and cyclophosphamide or combinations thereof.

Claim 33 (currently amended): A method for enhancing efficacy of a chemotherapeutic agent for a cancer cell, said method comprising administering systemically to a subject in need thereof an effective amount of a composition consisting essentially of hyaluronan and said chemotherapeutic agent, wherein the hyaluronan has a molecular weight between 400,000 and 900,000 modal molecular weight of 890,000 Da.

Claims 34-35 (canceled)

Claim 36 (currently amended): The method according to Claim [[34]] <u>33</u>, wherein the hyaluronan has a molecular weight of 890,000 Da.

Claim 38 (currently amended): The method according to Claim [[34]] <u>33</u>, wherein the chemotherapeutic agent is selected from the group consisting of methotrexate, paclitaxel, 5-fluorouracil and cyclophosphamide.

Claim 39 (currently amended): A method for overcoming acquired resistance of cancer cells to a chemotherapeutic agent, said method comprising administering <u>systemically</u> to a subject having said resistant cancer cells a hyaluronan and said chemotherapeutic agent, wherein the hyaluronan has a <u>molecular weight between 400,000 and 900,000 modal molecular weight of 890,000</u> Da.

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Claims 40-41 (canceled)

Claim 42 (currently amended): The method according to Claim [[40]] <u>39</u>, wherein the hyaluronan has a molecular weight of 890,000 Da.

Claim 43 (canceled)

Claim 44 (currently amended): The method according to Claim [[40]] <u>39</u>, wherein the chemotherapeutic agent is selected from the group consisting of methotrexate, paclitaxel, 5-fluorouracil and cyclophosphamide.

Claim 45 (currently amended): A pharmaceutical composition <u>formulated for systemic</u> <u>administration</u> consisting essentially of [[a]] <u>an anticancer</u> chemotherapeutic agent and hyaluronan, wherein the hyaluronan has a <u>molecular weight between 400,000 and 900,000 modal molecular weight of 890,000 Da.</u>

Claims 46-47 (canceled)

Claim 48 (currently amended): The pharmaceutical composition of Claim [[46]] <u>45</u>, wherein the hyaluronan has a molecular weight of 890,000 Da.

Claim 49 (canceled)

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Claim 50 (currently amended): The pharmaceutical composition of Claim [[46]] <u>45</u>, wherein the chemotherapeutic agent is selected from the group consisting of methotrexate, paclitaxel, 5-fluorouracil and cyclophosphamide.

Claim 51 (currently amended): A pharmaceutical composition <u>formulated for systemic administration</u> comprising [[a]] <u>an anticancer chemotherapeutic agent and hyaluronan having molecular weight of modal molecular weight of 890,000 Da.</u>

Claim 52 (previously presented): The pharmaceutical composition of Claim 51, wherein the hyaluronan has molecular weight 890,000 Da.

Claims 53-56 (canceled)

Claim 57 (new): The method of claim 27, wherein the hyaluronan has a polydispersity of 1.78.

Claim 58 (new): The method of claim 33, wherein the hyaluronan has a polydispersity of 1.78.

Claim 59 (new): The method of claim 27, wherein the hyaluronan and the chemotherapeutic agent are administered intravenously.

Claim 60 (new): The method of claim 33, wherein the hyaluronan and the chemotherapeutic agent are administered intravenously.

Claim 61 (new): The pharmaceutical composition of claim 45, wherein the hyaluronan has a polydispersity of 1.78.

Claim 62 (new): The pharmaceutical composition of claim 45, wherein the hyaluronan and the chemotherapeutic agent are formulated for intravenous administration.